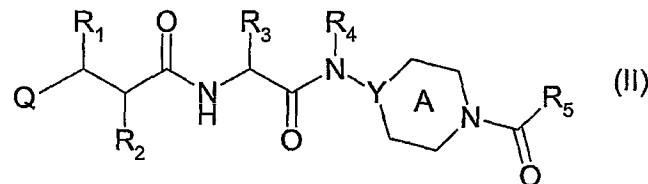


Correction of Listing of Claims for Preliminary Amendment**Listing of Claims:**

1. (Currently Amended) A compound of formula (II), or a pharmaceutical or veterinarily acceptable salt, hydrate or solvate thereof



wherein

Q represents a radical of formula -N(OH)CH(=O) or formula -C(=O)NH(OH);

R₁ represents hydrogen, methyl or trifluoromethyl, or, except when Z is a radical of formula -N(OH)CH(=O), a hydroxy, halo or amino group;

R₂ represents a group R₁₀-(V)_n-(ALK)_m- wherein

R₁₀ represents hydrogen, or a C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, cycloalkyl, aryl, or heterocyclyl group, any of which may be unsubstituted or substituted by (C₁-C₆)alkyl, (C₁-C₆)alkoxy, hydroxy, mercapto, (C₁-C₆)alkylthio, amino, halo-(including fluoro, chlorine, bromine and iodine), trifluoromethyl, cyano, nitro, oxo, -COOH, -CONH₂, -COOR^A, -NHCOR^A, -CONHR^A, -NHR^A, -NR^AR^B, or -CONR^AR^B wherein R^A and R^B are independently a (C₁-C₆)alkyl group and

ALK represents a straight or branched divalent C₁-C₆ alkylene, C₂-C₆ alkenylene, or C₂-C₆ alkynylene radical, and may be interrupted by one or more non-adjacent -NH-, -O- or -S-linkages,

V represents -NH-, -O- or -S-, and

m and n are independently 0 or 1;

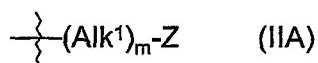
R₃ represents the side chain of a natural or non-natural alpha amino acid;

R₄ represents hydrogen or C₁-C₃ alkyl;

Y represents N or CH;

ring A is optionally substituted on one or more ring carbon atoms by C₁-C₃ alkyl, C₁-C₃ alkoxy, or halo; and

R₅ represents a group (IIA),



wherein

m is 0 or 1;

Alk¹ represents a divalent C₁-C₃ alkylene radical;

Z represents hydrogen or cycloalkyl, phenyl or heterocyclic which is optionally substituted by

(C₁-C₆)alkyl,

phenyl,

monocyclic 5 or 6-membered heterocyclic,

benzyl,

phenoxy, or (C₁-C₆)alkoxy,

phenylthio or (C₁-C₆)alkylthio, any of which is in turn optionally substituted by:

hydroxy or mercapto,

trifluoromethyl,

oxo,

nitro,

cyano (-CN),

bromo, chloro, fluoro, or iodo,

-COOH, or -COOR^A,

-CONH₂, -CONHR^A, or -CONR^AR^B

-COR^A, -SO₂R^A,

-NHCOR^A,

-NH₂, -NHR^A, or -NR^AR^B,

wherein R^A and R^B are independently a (C₁-C₆)alkyl group, or R^A and R^B taken together with the nitrogen atom to which they are attached form a 5- or 6-membered heterocyclic ring which may be substituted by -(C₁-C₃)alkyl(C₁-C₃)alkyl, hydroxy, or hydroxy(C₁-C₃)alkyl.

2. (Currently Amended) A compound as claimed in claim 1 wherein Z represents cycloalkyl, phenyl or monocyclic-heterocyclic, which is optionally substituted by (C₁C₆)alkyl, (C₂-C₆)alkenyl, or (C₂-C₆)alkynyl, phenyl, or halophenyl, trifluoromethyl, monocyclic 5 or 6-memberedhetrocyclic, benzyl, or halophenylmethyl,

hydroxy, phenoxy, (C₁-C₆)alkoxy, or hydroxy(C₁-C₆)alkyl,

mercapto, (C₁-C₆)alkylthio or mercapto(C₁-C₆)alkyl,

oxo,

nitro,

cyno (-CN),

bromo, chloro, fluoro, or iodo,

-COOH, or -COOR^A,

-CONH₂, -CONHR^A, or -CONR^AR^B

-COR^A, -SO₂R^A,

-NHCOR^A,

-NH₂, -NHR^A, or -NR^AR^B,

wherein R^A and R^B are independently a (C₁-C₆)alkyl group, or R^A and R^B taken together with the nitrogen atom to which they are attached form a 5- or 6-membered heterocyclic ring which may be substituted by (C₁-C₃)alkyl(C₁-C₃)alkyl, hydroxy, or hydroxy(C₁-C₃)alkyl.

3. (Currently amended) A compound as claimed in claim 1 or claim 2 wherein R₁ is hydrogen.

4. (Currently amended) A compound as claimed in ~~any of the preceding claims~~ claim 1 wherein R₂ is (C₁-C₆)alkyl-, cycloalkyl(C₁-C₆)alkyl-, (C₁-C₃)alkyl-S-(C₁-C₃)alkyl-, or (C₁-C₃)alkyl-O-(C₁-C₃)alkyl-.

5. (Currently amended) A compound as claimed in ~~any of claims 1 to 3~~ claim 1 wherein R₂ is n-propyl, n-butyl, n-pentyl, cyclopentylmethyl, cyclopentylethyl, cyclohexylmethyl or cyclohexylethyl.

6. (Currently amended) A compound as claimed in ~~any of the preceding claims~~ claim 1 wherein R₃ is

the characterising group of a natural α amino acid, ~~for example benzyl~~, or 4-methoxyphenylmethyl, in which any functional group may be protected, any amino group may be acylated and any carboxyl group present may be amidated; or a group -[Alk]_nR₉ where Alk is a (C₁-C₆)alkylene or (C₂-C₆) alkenylene group optionally interrupted by one or more -O-, or -S- atoms or -N(R₁₂)- groups [where R₁₂ is a hydrogen atom or a (C₁-C₆)alkyl group], n is 0 or 1, and R₉ is hydrogen or an optionally substituted phenyl, aryl, heterocyclyl, cycloalkyl or cycloalkenyl group or (only when n is 1) R₉ may additionally be hydroxy, mercapto, (C₁-C₆)alkylthio, amino, halo, trifluoromethyl, nitro, -COOH, -CONH₂, -COOR^A, -NHCOR^A, -CONHR^A, -NHR^A, -NR^AR^B, or -CONR^AR^B wherein R^A and R^B are independently a (C₁-C₆)alkyl group; or a benzyl group substituted in the phenyl ring by a group of formula -OCH₂COR₈ where R₈ is hydroxyl, amino, (C₁-C₆)alkoxy, phenyl(C₁-C₆)alkoxy, (C₁-C₆)alkylamino, di((C₁-C₆)alkyl)amino, phenyl(C₁-C₆)alkylamino; or a heterocyclic (C₁-C₆)alkyl group, either being unsubstituted or mono- or disubstituted in the heterocyclic ring with halo, nitro, carboxy, (C₁-C₆)alkoxy, cyano, (C₁-C₆)alkanoyl,

trifluoromethyl(C₁-C₆)alkyl, hydroxy, formyl, amino, (C₁-C₆)alkylamino, di-(C₁-C₆)alkylamino, mercapto, (C₁-C₆)alkylthio, hydroxy(C₁-C₆)alkyl, mercapto(C₁-C₆)alkyl or (C₁-C₆)alkylphenylmethyl; or

a group-CR_aR_bR_c in which:

each of R_a, R_b and R_c is independently hydrogen, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, phenyl(C₁-C₆)alkyl, (C₃-C₈)cycloalkyl; or

R_c is hydrogen and R_a and R_b are independently phenyl or heteroaryl such as pyridyl; or

R_c is hydrogen, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, phenyl(C₁-C₆)alkyl, or (C₃-C₈)cycloalkyl, and R_a and R_b together with the carbon atom to which they are attached form a 3 to 8 membered cycloalkyl or a 5-to 6-membered heterocyclic ring; or

R_a, R_b and R_c together with the carbon atom to which they are attached form a tricyclic ring (for example adamantly); or

R_a and R_b are each independently (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, phenyl(C₁-C₆)alkyl, or a group as defined for R_c below other than hydrogen, or R_a and R_b together with the carbon atom to which they are attached form a cycloalkyl or heterocyclic ring, and R is hydrogen, -OH, -SH, halogen, -CN, -CO₂H, (C₁-C₄)perfluoroalkyl, -CH₂OH, -CO₂(C₁-C₆)alkyl, -O(C₁-C₆)alkyl, -O(C₂-C₆)alkenyl, -S(C₁-C₆)alkyl, -SO(C₁-C₆)alkyl, -SO₂(C₁-C₆)alkyl, -SO₂(C₁-C₆)alkyl, -S(C₂-C₆)alkenyl, -SO(C₂-C₆)alkenyl, -SO₂(C₂-C₆)alkenyl or a group -Q-W wherein Q represents a bond or -O-, -S-, -SO- or -SO₂- and W represents a phenyl, phenylalkyl, (C₃-C₈)

cycloalkyl, (C₃-C₈) cycloalkylalkyl, (C₄-C₈) cycloalkenyl, (C₄-C₈) cycloalkenylalkyl, heteroaryl or heteroarylalkyl group, which group W may optionally be substituted by one or more substituents independently selected from, hydroxyl, halogen, -CN, -CO₂H, -CO₂(C₁-C₆)alkyl, -CONH₂, -CONH(C₁-C₆)alkyl alkyl, -CONH(C₁-C₆alkyl)₂, -CHO, -CH₂OH, (C₁-C₄)perfluoroalkyl, -O(C₁-C₆)alkyl, -S(C₁-C₆)alkyl, -SO(C₁-C₆)alkyl, -SO₂(C₁-C₆)alkyl, -NO₂, -NH₂, -NH(C₁-C₆)alkyl, -N((C₁-C₆)alkyl)₂, -NHCO(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, (C₃-C₈)cycloalkyl, (C₄-C₈)cycloalkenyl, phenyl or benzyl.

7. (Currently amended) A compound as claimed in ~~any of claims 1 to 6~~ claim 1 wherein R₃ is methyl, ethyl, n-propyl, n-butyl, benzyl, 4-chlorobenzyl, 4-hydroxybenzyl, phenyl, cyclohexyl, cyclohexylmethyl, pyridin-3-ylmethyl, tert-butoxymethyl, naphthylmethyl, iso-butyl, sec-butyl, tert-butyl, 1-benzylthio-1-methylethyl, 1-methylthio-1-methylethyl, 1- mercapto-1-methylethyl, 1-methoxy-1-methylethyl, 1-hydroxy-1-methylethyl, 1-fluoro- 1-methylethyl, hydroxymethyl, 2-hydroxethyl, 2-carboxyethyl, 2-methylcarbamoylethyl, 2-carbamoylethyl, or 4-aminobutyl.

8. (Currently amended) A compound as claimed in ~~any of claims 1 to 6~~ claim 1 wherein R₃ is tert-butyl, isobutyl, benzyl, isopropyl or methyl.

9. (Currently amended) A compound as claimed in ~~any of the preceding claims~~ claim 1 wherein R₄ is methyl.

10. (Currently amended) A compound, ~~method, use or composition as claimed in any of the preceding claims as claimed in claim 1~~ wherein in the group R₅, m is 1, and Alk¹ is -(CH₂)- or -(CH₂CH₂)-.
11. (Currently amended) A compound as claimed in ~~any of the preceding claims~~ claim 1 wherein, in the group R₅, Z is a phenyl, pyridyl, thienyl, furanyl, pyranyl, pyrolyl, diazolyl, triazolyl, thiazolyl, thiadiazolyl, oxazolyl, ozadiazolyl, indolyl, benzisozazolyl, benzthiazolyl or imidazothiazolyl ring, optionally substituted as specified in claim 1 ~~or claim 2~~.
12. (Original) A compound as claimed in claim 11 wherein the ring Z is unsubstituted or substituted by methyl, methoxy, ethoxy, methoxymethyl, ethylthio, chloro, bromo, hydroxy, nitro, phenyl, 2- or 4-nitrophenyl, dimethylamino, dimethylaminophenyl, methylsulphonyl, dimethylaminosulphonyl, 3-pyridyl or 2-pyrazin-2-yl.
12. (Canceled).
13. (Currently amended) A compound as claimed in ~~claim 1 or claim 2~~ wherein the compound is one specifically named and/or exemplified herein, or is the hydroxamate (Q represents a radical of formula -C(=O)NH(OH)) analogue thereof.
14. (Currently amended) A method for the treatment of bacterial infections in humans and

non-human mammals, which comprises administering to a subject suffering such infection an antibacterially effective dose of a compound as claimed in ~~any of claims 1 to 13~~claim 1.

15. (Currently amended) A method for the treatment of bacterial contamination by applying an antibacterially effective amount of a compound as claimed in ~~any of claims 1 to 13~~claim 1 to the site of contamination.

16. (Currently amended) The use of a compound as claimed in ~~any of claims 1 to 13~~claim 1 in the manufacture of an antibacterial composition.

17. (Currently amended) A pharmaceutical or veterinary composition comprising a compound as claimed in ~~any of claims 1 to 13~~claim 1 together with a pharmaceutical or veterinarily acceptable carrier.

18. (New) A compound as claimed in claim 1 wherein, in the group R₅, Z is a cyclopentyl, cyclohexyl, phenyl, morpholinyl, pyrimidin-2-yl, 1,2,3-thiadiazol-5-yl, 1,4-thiazol-5-yl, benzofuran-2-yl, 2- or 3-furanyl, 2- or 3-thienyl, 2- or 3-pyranyl, 2-, 3- or 4-pyrrolyl, 3-, 4- or 5-pyrazolyl, 3-, 4- or 5-isoxazolyl, or 2-, 3- or 4-pyridyl ring any of which may optionally be substituted by hydroxy, methoxy, ethoxy, mercapto, methylthio, ethylthio, methyl, ethyl, trifluoromethyl, fluoro, chloro, amino, methylamino, or dimethylamino.